EXHIBIT 3

UNITED STATES DISTRICT COURT DISTRICT OF MASSACHUSETTS

THE UNITED STATES OF AMERICA, ex rel. JULIE LONG,

Plaintiffs,

Civil Action No. 16-CV-12182-FDS

v.

JANSSEN BIOTECH, INC.,

Defendant.

DEFENDANT JANSSEN BIOTECH, INC.'S SUPPLEMENTAL INITIAL DISCLOSURES UNDER FEDERAL RULE OF CIVIL PROCEDURE 26(a)(1)

Pursuant to Rule 26(a)(1) of the Federal Rules of Civil Procedure, Defendant Janssen Biotech, Inc. ("Janssen" or "Defendant") provides the following supplemental initial disclosures based upon information reasonably available to it as of March 23, 2023.

The information provided in these disclosures related to Defendant's defenses is limited to the defenses that Defendant may assert in connection with the claims that Relator Julie Long ("Long" or "Relator") has asserted against Defendant. The information provided in these disclosures is also limited in accordance with the Court's decision that discovery be focused in the first instance on claims arising from support services provided by Long to the few physician practice accounts as alleged with any specificity in the Second Amended Complaint ("SAC"). Consistent with Rule 26(a), Defendant makes these disclosures to "accelerate the exchange of basic information about the case," *see* Notes of Advisory Committee on Rules—1993

Amendment, including by identifying the individuals and categories of documents that Janssen may use to support its claims and defenses. *See* Fed. R. Civ. P. 26(a)(1)(A)(i)–(ii).

By making these disclosures, Defendant does not concede that the prospective testimony of any individual identified herein, any document or category of documents identified herein, or any other information herein is relevant, discoverable, non-privileged, or admissible. In accordance with Rule 26(e), Defendant reserves the right to supplement or correct the disclosures set forth herein "in a timely manner" as necessary or appropriate. *See* Fed. R. Civ. P. 26(e)(1)(A).

A. Individuals Likely to Have Discoverable Information (Fed. R. Civ. P. 26(a)(1)(A)(i))

The following is a list of individuals that Defendant at the present time believes may have information that Defendant may use to support its defenses, excluding individuals with information that would be used solely for impeachment purposes. This list supplements and amends Defendant's disclosure concerning individuals likely to have discoverable information produced on December 22, 2020. For more information about individuals who Defendant believes might have generally discoverable information, Defendant directs you to the individuals identified as document custodians in this matter, and the individuals identified in Defendant's verified written responses to Relator's 30(b)(6) notice, which is attached here as Exhibit A.

Defendant does not consent to communications by Long or her counsel with any of the individuals identified below or with any of Defendant's other current or former employees as to matters related to this case. All such individuals may be contacted through Defendant's undersigned counsel.

Name	Subject(s)
Joseph Braunreuther	Healthcare compliance policies
Deputy General Counsel (prior)	and practices; disclosures of IOI
	Support Services ¹ to United

¹ The term IOI Support Services encompasses any in-office infusion-support program that Relator has identified in the SAC or her responses to Defendant's Interrogatory No. 2.

	States; Average Wholesale Price
	multidistrict litigation
Tom Cornely	Healthcare Compliance Policies;
HCC Officer, Immunology	Review and approval of IOI
Tiec officer, minimiology	Support Services
John Franz	Healthcare Compliance policies
	± ±
Healthcare Compliance Officer	and practices; Review and
Ken Gillmer	approval of IOI Support Services
	Remicade and Simponi ARIA
Director, Site of Care	marketing and IOI Support
F1 10 '1	Services; ABS responsibilities
Edmund Greenidge	Healthcare Compliance policies
Director, Health Care Compliance (prior)	and practices; Review and
D. G. 100 I	approval of IOI Support Services
Dana Griffith	Remicade and Simponi ARIA
Executive Area Business Specialist	marketing and IOI Support
7.4. 77.00	Services; ABS responsibilities
John Hoffman	Healthcare Compliance policies
Healthcare Compliance Officer	and practices; Review and
	approval of IOI Support Services;
	Average Wholesale Price
	multidistrict litigation
Marti Heckman	Remicade and Simponi ARIA
National Sales Director, Site of Care/ Immunology (prior)	marketing and IOI Support
	Services; ABS responsibilities
Kendra Heusinkveld	Remicade and Simponi ARIA
Product Manager, Immunology	marketing and IOI Support
	Services; ABS responsibilities
Freddy Jimenez	Healthcare compliance policies
Assistant General Counsel (prior)	and practices; Review and
	approval of IOI Support Services;
	disclosures of IOI Support
	Services to United States
James Knepp	Remicade and Simponi ARIA
Product Director, Site of Care Marketing & Head of	marketing and IOI Support
Communications Strategy for Site of Care	Services; ABS responsibilities
Julie Long	Remicade and Simponi ARIA
Area Business Specialist (prior)	marketing and IOI Support
1 4 /	Services; ABS responsibilities
Thao Marzullo	Remicade and Simponi ARIA
Product Manager, Site of Care	marketing and IOI Support
,	Services; ABS responsibilities
Maripat Rhood	Healthcare Compliance Policies;
Associate Director, Healthcare Compliance	Review and approval of IOI
1222 1 and 2 il control compilation	Support Services
	Support Services

Michael Schoeck	Healthcare Compliance Policies;
Director, Healthcare Compliance and Privacy	Review and approval of IOI
	Support Services
Scott Shelhamer	Remicade and Simponi ARIA
Director, U.S. Immunology Key Stakeholders	marketing and IOI Support
	Services; ABS responsibilities
Brian Smith	Remicade and Simponi ARIA
Director of Marketing, Immunology	marketing and IOI Support
	Services; Healthcare Compliance
	policies and practices; Review
	and approval of IOI Support
	Services
Karen Trahan	Remicade and Simponi ARIA
Regional Business Director (prior)	marketing and IOI Support
	Services; ABS responsibilities
Paul Wickmann	Remicade and Simponi ARIA
Regional Business Manager, Mid-Atlantic Region (prior)	marketing and IOI Support
	Services; ABS responsibilities
Michael Wolfe	Remicade and Simponi ARIA
Product Director, Site of Care Marketing	marketing and IOI Support
(prior)	Services; ABS responsibilities
Christopher Zalesky	Healthcare Compliance Policies;
Vice President Global Policy & Guidance (prior)	Review and approval of IOI
	Support Services; disclosures of
	IOI Support Services to United
	States

Discovery remains ongoing and Defendant may discover additional individuals with information that may be used to support its defenses. In particular, Defendant notes that it recently agreed to review and produce non-privileged responsive documents from more than 25 of the individuals disclosed in Exhibit A. Consistent with Fed. R. Civ. P. 26, if during the course of discovery, Defendant learns of additional witnesses, it reserves the right to supplement the list set forth above as necessary or appropriate. Defendant further reserves the right to identify and call as witnesses at trial or otherwise additional persons who, during the course of discovery and investigation relating to this case, Defendant learns may have knowledge of relevant matters. Furthermore, for the avoidance of doubt, though Janssen has identified two individuals who served in legal roles at the company in the above disclosure of individuals likely to have

discoverable information, it does not list these individuals because it intends to rely on privileged legal advice they provided about the IOI Support Services. Instead, as specified above, they are listed as potential fact witnesses regarding healthcare compliance policies and practices and disclosures of information to the government through previous investigations and litigation. As previously stated, Janssen has not asserted and does not intend to assert the advice of counsel defense. Defendant reserves the right to object to the relevance or admissibility of any testimony of the individuals listed above if offered by Long or any other party.

B. Description of Relevant Documents (Fed. R. Civ. P. 26(a)(1)(A)(ii))

The following is a description, by category, of documents, data compilations, and tangible things known to Defendant and in its possession, custody, or control that may be used to support its defenses to Long's claims, excluding items that may be used solely for impeachment.

- Slide decks, brochures, presentations, and related materials for the IOI Support Services that Long and other Area Business Specialists ("ABSs") presented to the physician practice accounts identified in the SAC;
- Documents related to communications between Long, or any other ABS, and the physician practice accounts identified in the SAC;
- Agreements with any third-party vendors that assisted Long and other ABSs in providing support services to the physician practice accounts identified in the SAC;
- Information about sales of Remicade, Simponi ARIA, and competitor medications to the physician practice accounts identified in the SAC;
- Documents relating to training, direction, and instruction given to Long and other ABSs concerning their role, responsibilities, performance expectations, interactions with physician practices, and presentation of the IOI Support Services provided to the physician practice accounts identified in the SAC;
- Documents related to Janssen's healthcare compliance policies and procedures, including training on compliance policies, implementation guidance, and reporting and adjudication of potential compliance violations;
- Non-privileged documents relating to the submission of the IOI Support Services for Promotional Review Committee approval;

- Non-privileged documents relating to the review and approval of the IOI Support Services by the Promotional Review Committee;
- Documents reflecting publicly available information about opening and managing inoffice-infusion suites for patients;
- Guidance and advisory opinions published by the Office of the Inspector General for the Department of Health and Human Services, or any other government entity, related to healthcare compliance by pharmaceutical manufacturers, the Anti-Kickback statute, and the provision of information or services to healthcare providers;
- Data from the Centers for Medicare & Medicaid Services ("CMS") related to prescriptions and infusions of Remicade, Simponi Aria, and other competing therapies;²
- Documents related to the development, creation, use, promotion, and evaluation of the IOI Support Services;
- Documents relating to disclosure of the IOI Support Services to the United States, including in response to requests related to the document request issued in 2003 by the United States Department of Justice, and disclosed through litigation involving the United States;
- Documents relating to healthcare industry guidance and practice concerning the provision of information to physician practices and other healthcare providers as part of sales and marketing;
- Documents relating to the business and marketing purposes of the IOI Support Services;
- Documents, including exhibits, trial transcripts, and filings from prior litigation involving issues related to the provision of IOI Support Services.

Defendant's categorical descriptions of the documents it may rely on to support its claims and defenses is made without prejudice to its ability to, consistent with Fed. R. Civ. P. 26(a)(3), identify the specific documents it may rely on at trial when appropriate. *See* Notes of Advisory Committee on Rules—1993 Amendment (explaining that "unlike subdivision (a)(3)(C), an itemized listing of each exhibit is not required [under 26(a)(1)(A)(ii)]," and noting that instead,

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² For clarity, the only relevant CMS data in Janssen's possession about which it is currently aware, is the CMS data produced to it by Relator in this litigation.

under 26(a)(1)(A)(ii) "the disclosure should describe and categorize, to the extent identified during the initial investigation, the nature and location of potentially relevant documents and records "). Defendant may discover additional categories of documents that may be used to support its defenses during the course of discovery and, consistent with Fed. R. Civ. P. 26, reserves the right to supplement the list set forth above as necessary or appropriate. Defendant reserves the right to object to the relevance or admissibility of any of the documents identified above if offered by Long or any other party for any purpose.

C. Computation of Damages (Fed. R. Civ. P. 26(a)(1)(A)(iii))

Defendant does not, at this time, seek damages from Long as to which disclosure is required pursuant to Rule 26(a)(1)(A)(iii) of the Federal Rules of Civil Procedure, although Defendant may seek costs and expenses incurred in this action, including attorney's fees.

D. Insurance Agreements (Fed. R. Civ. P. 26(a)(1)(A)(iv))

Defendant is currently unaware of any policy or insurance under which any person or entity may be liable to satisfy all or part of a possible judgment in this action or to indemnify or reimburse for payments made to satisfy the judgment.

* * *

Nothing in Defendant's initial disclosure of witnesses, documents, or other information shall constitute an admission, concession, or waiver with respect to any issue of fact or law, or any claim, defense, or privilege including (without limitation) the following: any claim or defense as to the sufficiency of Relator's SAC; the relevance, discoverability, or admissibility of any of the information set forth herein; any applicable privilege or immunity, including the attorney-client privilege, the work product privilege, or any other privilege or immunity; and the right to object to discovery requests that are not relevant, unduly burdensome, not proportional, and/or not reasonably calculated to lead to the discovery of admissible evidence.

Defendant reserves the right to supplement or modify these disclosures as necessary or appropriate.

Dated: March 23, 2023 s/Jason C. Raofield

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CERTIFICATE OF SERVICE

The undersigned hereby certifies that a true and correct copy of the foregoing document has been served by electronic mail on March 23, 2023, to the following counsel of record:

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EXHIBIT A

UNITED STATES DISTRICT COURT DISTRICT OF MASSACHUSETTS

THE UNITED STATES OF AMERICA, ex rel. JULIE LONG,

Plaintiffs,

Civil Action No. 16-CV-12182-FDS

v.

JANSSEN BIOTECH, INC.,

Defendant.

<u>DEFENDANT JANSSEN BIOTECH, INC.'S AMENDED WRITTEN RESPONSES TO RELATOR'S NOVEMBER 4, 2022 30(B)(6) DEPOSITION NOTICE</u>

Pursuant to the Court's March 9, 2023 Order, Defendant Janssen Biotech, Inc. ("Janssen" or "Defendant") provides the following amended objections and responses to Relator's Notice of Deposition pursuant to Federal Rule of Civil Procedure 30(b)(6) (the "30(b)(6) Notice"), dated November 4, 2022.

GENERAL OBJECTIONS

- 1. Janssen objects to each Topic set forth in the Notice to the extent that it calls for information protected from disclosure by the attorney-client privilege, the protection afforded work product, or any other applicable privilege or protection. As previously stated, Janssen has not asserted and does not intend to assert the advice of counsel defense. For avoidance of doubt, by providing the requested factual information regarding the identities of the individuals who provided privileged legal advice, Janssen is not asserting the advice of counsel defense and has not waived any applicable privilege.
- 2. Janssen objects to each Topic set forth in the Notice to the extent it is vague and ambiguous, or fails to state with "reasonable particularity the matters for examination" as required by Rule 30(b)(6) of the Federal Rules of Civil Procedure.

- 3. Janssen objects to each Topic to the extent that it seeks information not relevant to any claim, counterclaim, or defense pled by any party in this action.
- 4. Janssen objects to each Topic to the extent that it is overly broad and unduly burdensome and not proportional to the needs of the case, and thus exceeds the scope of discoverable matters under Fed. R. Civ. P. 26.
- 5. Janssen objects to each Topic set forth in the Notice to the extent that it calls for information that is confidential or proprietary to Janssen. To the extent that Janssen provides such information, it will do so pursuant to the terms of the Protective Order entered in this case on March 11, 2021.
- 6. Janssen objects to each Topic set forth in the Notice to the extent it purports to impose any obligation on Janssen to locate, obtain, and/or provide testimony or documents regarding information not in the possession, custody, or control of Janssen.
- 7. Janssen objects to each Topic set forth in the Notice to the extent it seeks to impose obligations or requirements on Janssen which are greater than or different from those imposed by the Federal Rules of Civil Procedure and/or any other applicable law, rule, or regulation.
- 8. Janssen objects to any implications and to any explicit or implicit characterization of facts, events, circumstances, or issues set forth in the Notice. Any response by Janssen is not intended to indicate that Janssen agrees with any implications or any explicit or implicit characterization of facts, events, circumstances, or issues in the Notice, or that such implications or characterizations are relevant to this action.
- 9. Janssen objects to each Topic set forth in the Notice to the extent it seeks information outside of the scope authorized by the Court. Specifically, Janssen objects to each

Topic to the extent it seeks information outside the scope of the phased discovery authorized by the Court on December 14, 2020, and in subsequent orders by Chief Magistrate Judge Kelley, to the extent it seeks information that is not related to support services provided by Relator during her employment as an ABS at Janssen to the physician practices specifically alleged in the Second Amended Complaint.

10. Janssen's responses are made without waiving or intending to waive in any way (a) any objections as to competency, relevancy, materiality, privilege and/or admissibility, or subject matter thereof, in any subsequent proceeding in this or any other action; (b) the right to object on any ground to the use of these responses, or the subject matter thereof, in any subsequent proceeding in this or any other action; (c) the right to object to a demand for further responses to these or any other discovery involving or related to the subject matter of these topics; and (d) the right to object on any ground to these or any other or future discovery responses.

SPECIFIC OBJECTIONS AND RESPONSES

TOPIC NO. 1: The persons (excluding the PRC) that had significant involvement in the development and/or approval of the strategy or plan to provide and/or continue to provide any or all of the IOI Support Services (including the specific IOI Support Service(s) in which each such person had the significant involvement, the approximate date(s) of the significant involvement, the person's last known address and phone number, and, if an individual, the position they held at the time of their significant involvement).

<u>Relevant time period</u>: From when the strategy of providing the IOI Support Services was devised until February 19, 2016.

RESPONSE TO TOPIC NO. 1:

Janssen objects to this Topic to the extent that it is overly broad and unduly burdensome and not proportional to the needs of the case, and thus exceeds the scope of discoverable matters under Federal Rule of Civil Procedure 26.

Janssen further objects to this Topic to the extent it is cumulative and seeks information that is duplicative of what is contained in the documents or written discovery that Janssen has already produced.

Janssen further objects to this Topic on the ground that it is vague, ambiguous, and overbroad and does not describe with reasonable particularity the matters on which examination is requested as required by Fed. R. Civ. P. 30(b)(6). For example, Janssen objects to the Topic's use of the terms "development," "approval," "strategy," "plan," and "significant involvement," as those are not defined terms. "Significant involvement," in particular, is an inherently subjective term, and reasonable minds could differ as to whether a particular person was "significantly" involved in any given activity. This response is based on Janssen's good faith understanding, and it seeks to address Relator's concerns about providing a response that is overinclusive or under-inclusive.

Further, pursuant to the Court's March 9, 2023 order, Janssen's response will "only be with regard to the services at issue in this case, not all services." *See* ECF No. 375 at 18.

Accordingly, Janssen's response to Topic No. 1 will be with respect to the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2. *See* Relator's 3d Supplemental Objections and Responses to Janssen's Rog. No. 2 (January 27, 2023).

Although Topic No. 1 requests information through February 16, 2016, Janssen has provided responsive information through February 2020.

Subject to and without waiving the foregoing objections, Janssen responds as follows:

The individuals with significant involvement in the initial strategy to provide educational support to physicians (which began in approximately 1998 when Remicade was first approved) and the development of the initial educational materials related to IOI (the "Practice

Management Program") were Mike Ziskind, Brian Fitzpatrick, Julie McHugh, and Chris Zalesky.

The development and/or approval of the strategy or plan to provide and/or continue to provide the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2 was a function of certain members of the Site of Care ("SOC") team. The SOC team was responsible for developing educational materials for use by the Area Business Specialists ("ABS") with all sites of care for Remicade and Simponi Aria, such as: (1) Rheumatologists, Gastroenterologists, and Dermatologists; and (2) "Alternative Sites of Care," including Hospital Outpatient Departments and Infusion Therapy Providers, which included Ambulatory Infusion Centers and Home Infusion Providers.

Given the number of functions served by the SOC team, as a general matter, the individuals significantly involved in the development and/or approval of the strategy or plan to provide and/or continue to provide the IOI Support Services were the Product Director and Product Manager responsible for the IOI educational materials used by the ABSs. Generally, either the SOC Product Director or the Product Manager responsible for the IOI educational materials was the individual identified as the "Project Owner" on the PRC forms that have been produced in this matter. Further, as a general matter, Group Product Directors, Directors, and Vice Presidents (as identified on the organizational charts produced in this litigation) were not significantly involved in the development and/or approval of the strategy or plan to provide and/or continue to provide the IOI Support Services, given their broad range of responsibilities.

The SOC team has been housed in different groups of the Immunology Division.

Specifically, in 2003, the SOC team was part of the Strategic Customer Relations Group. *See*JANSSENBIO-042-00000024-37 at JANSSENBIO-042-00000035. In 2006, the SOC team was

part of the Strategic Business Franchise. *See* JANSSENBIO-042-00000093-103 at 101. In 2007, the SOC team was its own group within the Immunology Division. *See* JANSSENBIO-042-00000104-117 at 112. Between 2008 and the end of 2009, the SOC team was combined with Academic Marketing (the team responsible for marketing related to educational institutions, such as universities). *See* JANSSENBIO-042-00000129. Between 2010 and 2011, the SOC team was combined with Payer Marketing (the team responsible for marketing related to insurance companies). JANSSENBIO-042-00000132; JANSSENBIO-042-00000125. In the Fall of 2011, the SOC team was divided between the Rheumatology Group and the Gastroenterology Group. *See* JANSSENBIO-042-00000136. In 2014, Janssen again reorganized, and the SOC team was placed in the Immunology Inc. Portfolio group. *See* JANSSENBIO-042-00000124. As of February 2020, the SOC team remained a part of this group.

The following list encompasses the individuals significantly involved in the development and/or approval of the strategy or plan to provide and/or continue to provide the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2:

Individual	Title	IOI Support Program(s)
John Arena	Product Manager, Site of Care (May 2009– June 2010); Product Director, Site of Care (July 2010–	Significantly involved in the SOC strategy and the development of IOI educational materials in use between
	approx. 2012)	May 2009 and 2012. Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services: Considerations for Proactive Practice
		Management

¹ In 2016, the Immunology Inc. Portfolio Group was renamed "Immunology Portfolio Solutions." *See* JANSSENBIO-042-00000119. In 2018, the Immunology Portfolio Solutions group was renamed "Immunology Biosimilar and Portfolio Strategy." *See* JANSSENBIO-047-00009633.

Individual	Title	IOI Support Program(s)
Janice Babia- Ramos	Manager, Health Economics & Reimbursement (Jan. 2003–Jan. 2005); Senior Manager, Site of Care (Feb. 2005–Mar. 2005); Associate Director, Site of Care (Mar. 2005–May 2005)	Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI Support Services, in use between January 2003 and May 2005. Listed as the Project Owner on relevant PRC forms, including for at least the following IOI educational materials:
Darah Biddle	Product Manager, Crohn's Disease (Aug. 2010–May 2012); Product Director, Institutional Marketing (May 2012–Apr. 2014)	Infusion Optimization Modeler (IOM) Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services:
		Infusion Therapy Services Provided in Converted ASC Space; IV Therapy: An Important Option for Your Patients (a/k/a Why IV?); Successful Implementation of a New Infusion Suite for Gastroenterology Practices; Successful Infusion Suite Management; Successful Infusion Suite Management for Gastroenterology
Michelle Carrigan	Product Director, Channel Marketing (Nov. 2019–Mar. 2022)	Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI Support Services, in use between November 2019 and February 2020.
Lawrence Conley	Product Manager, Site of Care/Institutional Marketing (Feb. 2013–Oct. 2013); Product Manager, Gastroenterology (Nov. 2013–Apr. 2016)	Significantly involved in the SOC strategy and the development of IOI educational materials in use between February 2013 and April 2016. Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services:
		Infusion Optimization Modeler (IOM);

Individual	Title	IOI Support Program(s)
		Infusion Services Review (iBiz); IV Therapy: An Important Option for Your Patients (a/k/a Why IV?)
Camille Dorsey	Product Manager, Remicade (April 2012– May 2015); Product Director (Mar. 2015–Nov. 2017)	Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services:
		Considerations for Proactive Practice Management; Successful Infusion Suite Management for Gastroenterology
Joseph Donahue	Product Manager, Site of Care (May 2006– June 2007)	Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI Support Services, in use between May 2006 and June 2007.
		Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services:
		Setting Up In-Office Infusions of Remicade
Maria Finlay	Product Manager, Immunology Marketing (Sept. 2010–Apr. 2013)	Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI Support Services, in use between September 2010 and April 2013. Listed as the Project Owner on
		relevant PRC forms, including for at least the following IOI Support Services:
		Raising the Infusion Suite Experience (RISE)
Ken Gillmer	Senior Product Manager, Site of Care Marketing (June 2006–Aug. 2007); Associate Director, Site of Care (Aug. 2007–Oct. 2008); Director, Site of Care Marketing (Oct. 2008–Oct. 2010)	Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI Support Services, in use between June 2006 and October 2010.

Individual	Title	IOI Support Program(s)
		Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services:
		Becoming an Alternative Site of Care; Considerations for Proactive Practice Management; Enhancing Patient Care and Access; Managing Biologics in the Physician Office; Practice Compliance for Remicade; Private Payer Contracting Considerations; Setting Up In-Office Infusions of Remicade
Kendra Heusinkveld	Product Manager, Emerging Stakeholders (Oct. 2015–May 2017)	Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI Support Services, in use between October 2015 and May 2017.
		Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services:
		Infusion Services Review (iBiz); Infusion Suite Scheduling & Staffing; Quality of Care in the Infusion Suite; In-Office Infusion Drug Procurement Models;
		Inventory and Supply Management; Billing and Coding for Infusions; ICD-10; Clinical Protocols and SOPs in the Infusion Suite;
		Payer Relationship Management; Infusion Referrals
Jim Knepp	Product Director, Site of Care (June 2012–October 2015)	Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI Support Services, in use between June 2012 and October 2015.

Individual	Title	IOI Support Program(s)
		Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services:
		IV Therapy: An Important Option for Your Patients (a/k/a Why IV?)
Hitu Malhotra	Product Manager, Remicade (Sept. 2010– Apr. 2012)	Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI Support Services, in use between September 2010 and April 2012.
		Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services:
		Considerations for Proactive Practice Management; Enhancing Infusion Efficiency; Infusion Optimization Modeler (IOM); Managing Biologics in the Physician Office; Practice Compliance for Remicade; Private Payer Contracting Considerations: Part 2; Successful Implementation of a New Infusion Suite
Thao Marzullo	Product Manager, Site of Care Marketing (July 2013–Aug. 2015)	Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI Support Services, in use between July 2013 and August 2015.
		Listed as the Project Owner on relevant PRC forms, including for at least the following IOI Support Services:
Randy McGonigal	Senior Director, Site of Care Marketing (Dec. 2002–Jan. 2010)	Payer Relationship Management Significantly involved in the SOC strategy and the development of IOI educational materials, including IOI

Individual	Title	IOI Support Program(s)
		Support Services, in use between
		December 2002 and January 2010.
Tim Michael	Product Director, Site of Care (May 2016–	Significantly involved in the SOC
	June 2019)	strategy and the development of IOI
		educational materials in use between
		May 2016 and June 2019.
		Listed as the Project Owner on
		relevant PRC forms, including for at
		least the following IOI Support
		Services:
		11 PM 25 0 15
		Hot Buttons and Practice Pearls -
		Payer Relationship Management;
		Hot Buttons and Practice Pearls -
		Inventory and Supply Management;
		Hot Buttons and Practice Pearls - In-
		Office Infusion Drug Procurement Models;
		Billing and Coding for Infusions;
		Considerations for Proactive Practice
		Management
Tom Nyairo	Associate Product Manager, Care Delivery	Significantly involved in the SOC
The second	Strategies (Aug. 2004-Nov. 2006)	strategy and the development of IOI
		educational materials, including IOI
		Support Services, in use between
		August 2004 and November 2006.
		Listed as the Project Owner on
		relevant PRC forms, including for at
		least the following IOI Support
		Services:
		Enhancing Patient Care and Access;
		Infusion Optimization Modeler
		(IOM);
		Infusion Therapy Services Provided in Converted ASC Space;
		Remicade Account Review
Bill Pillat	Product Manager, Site of Care (May 2017–	Significantly involved in the SOC
	Aug. 2019)	strategy and the development of IOI
	mis amis	educational materials, including IOI
		Support Services, in use between May
		2017 and August 2019.

Individual	Title	IOI Support Program(s)
		Listed as the Project Owner on
		relevant PRC forms, including for at
		least the following IOI Support
		Services:
		Infusion Services Review (iBiz);
		Quality of Care in the Infusion Suite;
		In-Office Infusion Drug Procurement
		Models;
		Infusion Suite Scheduling & Staffing;
		Specialty Drug Market Dynamics:
		Implications for Infusions;
		Inventory and Supply Management
Faiz Sadeq	Product Manager, Alternative Site of Care	Significantly involved in the SOC
	(Dec. 2010–July 2011);	strategy and the development of IOI
	Product Manager, Site of Care/Hospital	educational materials, including IOI
	(July 2011–Dec. 2012)	Support Services, in use between
		December 2010 and December 2012.
		Listed as the Project Owner on
		relevant PRC forms, including for at
		least the following IOI Support
		Services:
		Services.
		Checkpoints for Infusion Center
		Optimization;
		Managing Biologics in the Physician
		Office; Private Payer Contracting
		Considerations
Scott	Regional Business Director (Apr. 2015–	Significantly involved in the SOC
Shelhamer	Feb. 2020)	strategy and the development of IOI
		educational materials, including IOI
		Support Services, in use between
		April 2015 and February 2020.
Sandra	Product Manager (2002–2005)	Significantly involved in the SOC
Shpilberg		strategy and the development of IOI
		educational materials in use between
		Listed as the Project Owner on
		relevant PRC forms, including for at
		least the following IOI Support
		Services:
		Setting Up In-Office Infusions of
		Remicade;

Individual	Title	IOI Support Program(s)
		Private Payer Contracting
		Considerations for Remicade;
		Remicade Account Review (Physician
		Office Account Review for Remicade)
David Silver	Product Director, Remicade (Aug. 2008–	Listed as the Project Owner on
	Aug. 2011)	relevant PRC forms, including for at
		least the following IOI Support
		Services:
		F - 1 - 7 - 1 1 77 - 14
	Description District Property (2000 2015)	Emerging Trends in Healthcare
Karen Trahan	Regional Business Director (2000–2015)	Significantly involved in the SOC
		strategy and the development of IOI
		educational materials, including IOI
2		Support Services, in use between 2000 and 2015.
Michael Wolfe	Product Director, Marketing, Site of Care	Significantly involved in the SOC
	(Oct. 2010–Apr. 2012);	strategy and the development of IOI
	Product Director, Rheumatology	educational materials, including IOI
	Marketing (Apr. 2012–Apr. 2016)	Support Services, in use between
		October 2010 and April 2016.
		Listed as the Project Owner on
		relevant PRC forms, including for at
		least the following IOI Support Services:
		Services:
		Managing Biologics in the Physician
		Office; Successful Implementation of
		a New Infusion Suite
David Wright	Group Product Director, Stakeholder (Feb.	Significantly involved in the SOC
	2013-Dec. 2014);	strategy and the development of IOI
	Group Product Director, Rheumatology,	educational materials, including IOI
	IV/IOI (Dec. 2014–Mar. 2016)	Support Services, in use between
		February 2013 and March 2016.
Michael	Director, Public Payer Policy, Strategy &	Significantly involved in the SOC
Ziskind	Marketing (Centocor) (1998-2006)	strategy and the development of IOI
	20 Sept. 20	educational materials, including IOI
		Support Services, in use between 1998
		and 2006.
		Listed as the Project Owner on
		relevant PRC forms, including for at
		least the following IOI Support
		Services:

Individual	Title	IOI Support Program(s)
		Infusion Therapy Services Provided in
		ASC;
		Managing Biologics in the Physician's
		Office

<u>TOPIC NO. 2:</u> The persons (excluding the PRC) that had significant involvement in the decision to stop providing any or all of the IOI Support Services (including the specific IOI Support Service(s) in which each such person had the significant involvement, the approximate date(s) of the significant involvement, the person's last known address and phone number, and, if an individual, the position they held at the time of their significant involvement).

Relevant time period: From April 9, 2003 until February 19, 2020.

RESPONSE TO TOPIC NO. 2:

Janssen objects to this Topic to the extent that it is overly broad and unduly burdensome and not proportional to the needs of the case, and thus exceeds the scope of discoverable matters under Federal Rule of Civil Procedure 26.

Janssen further objects to this Topic to the extent it is cumulative and seeks information that is duplicative of what is contained in the documents or written discovery that Janssen has already produced.

Janssen further objects that this Topic on the ground that it is vague, ambiguous, and overbroad and does not describe with reasonable particularity the matters on which examination is requested as required by Fed. R. Civ. P. 30(b)(6). For example, Janssen objects to the Topic's use of the term "significant involvement," as that term is not defined. "Significant involvement," in particular, is an inherently subjective term, and reasonable minds could differ as to whether a particular person was "significantly" involved in any given activity. This response is based on Janssen's good faith understanding, and it seeks to address Relator's concerns about providing a response that is over-inclusive or under-inclusive.

Further, pursuant to the Court's March 9, 2023 order, Janssen's response will "only be with regard to the services at issue in this case, not all services." *See* ECF No. 375 at 18.

Accordingly, Janssen's response to Topic No. 2 will be with respect to the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2. *See* Relator's 3d Supplemental Objections and Responses to Janssen's Rog. No. 2 (January 27, 2023).

Subject to and without waiving the foregoing objections, Janssen responds as follows:

As a general matter, the individuals significantly involved in the decision to discontinue particular IOI Support Services were the Product Director and Product Manager responsible for the IOI educational materials used by the ABSs.

The following list encompasses the individuals significantly involved in in the decision to discontinue particular IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2:

Individual	Title	IOI Support Program(s)
John Arena	Product Manager, Site of Care (May 2009-	Significantly involved in the decision
	June 2010);	to stop providing any IOI Support
	Product Director, Site of Care (July 2010–approx. 2012)	Services between May 2009 and 2012.
	(D) Anisa 3.550	Programs that were discontinued
		between May 2009 and 2012 included:
		Managing Biologics in the Physician Office;
		Successful Infusion Site Management
		for Gastroenterology;
		Successful Implementation of a New
		Infusion Suite for Gastroenterology Practices;
		Successful Implementation of a New
		Infusion Suite;
		Private Payer Contracting
		Considerations – Part 1 and Part 2
		(a/k/a Private Payer Contracting
		Considerations for Therapy With
		Remicade);
		Enhancing Efficiency and Capacity;
		Considerations for Working With a
		Specialty Pharmacy;

Individual	Title	IOI Support Program(s)
		Becoming an Alternative Site of Care
		for Therapy with Remicade in Your
		Community
Janice Babia-	Manager, Health Economics &	Significantly involved in the decision
Ramos	Reimbursement (Jan. 2003-Jan. 2005);	to stop providing any IOI Support
	Senior Manager, Site of Care (Feb. 2005-	Services between January 2003 and
	Mar. 2005);	May 2005.
	Associate Director, Site of Care (Mar.	STANDARD CONTRACTOR AND STANDARD
	2005–May 2005)	Programs that were discontinued
	Table According to 10 Membrane Management of the Control of the Co	between January 2003 and May 2005
		included:
		5 (5 () () () () () () () () (
		Infusion Suite Designer
Michelle	Product Director, Channel Marketing	Significantly involved in the decision
Carrigan	(Nov. 2019–Mar. 2022)	to stop providing any IOI Support
	31 340	Services between November 2019 and
		February 2020.
Lawrence	Product Manager, Site of Care/Institutional	Significantly involved in the decision
Conley	Marketing (Feb. 2013–Oct. 2013);	to stop providing any IOI Support
3. In a 2.	Product Manager, Gastroenterology (Nov.	Services between February 2013 and
	2013–Apr. 2016)	April 2016.
	· **	<u> </u>
		Programs that were discontinued
		between January 2013 and April 2016
		included:
		Checkpoints for Infusion Center
		Optimization;
		Efficiency Checklist;
		Infusion Optimization Modeler (IOM);
		Setting Up In-Office Infusions of
		Remicade: Informational Resources
Joseph	Product Manager, Site of Care (May 2006–	Significantly involved in the decision
Donahue	June 2007)	to stop providing any IOI Support
	9100 - CUIZ	Services between May 2006 and June
		2007.
		Programs that were discontinued
		between May 2006 and June 2007
		included:
		Remicade Account Review (a/k/a
		Physician Office Account Review for
1		Remicade)

Individual	Title	IOI Support Program(s)
Ken Gillmer	Senior Product Manager, Site of Care Marketing (June 2006–Aug. 2007); Associate Director, Site of Care (Aug. 2007–Oct. 2008); Director, Site of Care Marketing (Oct. 2008–Oct. 2010)	Significantly involved in the decision to stop providing any IOI Support Services between June 2006 and October 2010. Programs that were discontinued
	2008–001. 2010)	between June 2006 and October 2010 included:
		Remicade Account Review (a/k/a Physician Office Account Review for Remicade)
Kendra Heusinkveld	Product Manager, Emerging Stakeholders (Oct. 2015–May 2017)	Significantly involved in the decision to stop providing any IOI Support Services between October 2015 and May 2017.
		Programs that were discontinued between October 2015 and May 2017 included:
		Considerations for Proactive Practice Management (a/k/a Current Considerations for Proactive Practice Management; Proactive Practice
		Management); Considerations for Standard Operating Procedures in the Infusion Suite; Electronic Health Records and Meaningful Use;
		Electronic Health Records and Meaningful Use 2 (a/k/a EHR/MU2); Payer Relationship Management; Practice Compliance for Remicade
Jim Knepp	Product Director, Site of Care (June 2012–October 2015)	Significantly involved in the decision to stop providing any IOI Support Services between June 2012 and October 2015.
		Programs that were discontinued between June 2012 and October 2015 included:
		Checkpoints for Infusion Center Optimization;

Individual	Title	IOI Support Program(s)
		Efficiency Checklist;
		Infusion Optimization Modeler (IOM);
		Infusion Therapy Services Provided in
		Converted ASC Space (a/k/a Infusion
		Services and Ambulatory Surgical
		Centers (ASCs) – Planning
		Considerations;
		ASC Space Reclassification for
		Infusion Therapy);
		Private Payer Contracting
		Considerations – Part 1 and Part 2
		(a/k/a Private Payer Contracting
		Considerations for Therapy With
		Remicade);
		Setting Up In-Office Infusions of
TT' 3 5 11	D 1 116 D 1 16 1 2012	Remicade: Informational Resources
Hitu Malhotra	Product Manager, Remicade (Sept. 2010–	Significantly involved in the decision
	Apr. 2012)	to stop providing any IOI Support
		Services between September 2010 and
		April 2012.
		Dro grows that were discontinued
		Programs that were discontinued
		between June September 2010 and
		April 2012 included:
		Becoming an Alternative Site of Care
		for Therapy with Remicade in Your
		Community;
		Considerations for Working With a
		Specialty Pharmacy;
		Enhancing Efficiency and Capacity;
		Private Payer Contracting
		Considerations – Part 1 and Part 2
		(a/k/a Private Payer Contracting
		Considerations for Therapy With
		Remicade)
Thao	Product Manager, Site of Care Marketing	Significantly involved in the decision
Marzullo	(July 2013–Aug. 2015)	to stop providing any IOI Support
		Services between July 2013 and August
		2015.
		Programs that were discontinued
		between June July 2013 and August
		2015 included:

Individual	Title	IOI Support Program(s)
		Efficiency Checklist;
		Infusion Optimization Modeler (IOM);
		Infusion Therapy Services Provided in
		Converted ASC Space (a/k/a Infusion
		Services and Ambulatory Surgical
		Centers (ASCs) – Planning
		Considerations; ASC Space
		Reclassification for Infusion Therapy);
		Setting Up In-Office Infusions of
		Remicade: Informational Resources
Randy	Senior Director, Site of Care Marketing	Significantly involved in the decision
McGonigal	(Dec. 2002–Jan. 2010)	to stop providing any IOI Support
		Services between December 2002 and
		January 2010.
		Programs that were discontinued
		between June December 2002 and
		January 2010 included:
		Infusion Suite Designer;
		Remicade Account Review (a/k/a
		Physician Office Account Review for
		Remicade)
Tim Michael	Product Director, Site of Care (May 2016–	Significantly involved in the decision
	June 2019)	to stop providing any IOI Support
		Services between May 2016 and June
		2019.
		Programs that were discontinued
		between June May 2016 and June 2019
		included:
		Dilling and Coding for Inferiors
		Billing and Coding for Infusions; Considerations for Proactive Practice
		Characteristics and the selection reached and discount and the contraction of the selection of the selection of
		Management (a/k/a Current Considerations for Proactive Practice
		Management; Proactive Practice
		Management);
		Electronic Health Records and
		Meaningful Use;
		Electronic Health Records and
		Meaningful Use 2 (a/k/a EHR/MU2);
		Emerging Trends in Healthcare;
		Infusion Services Review (iBiz);
		Infusion Suite Scheduling and Staffing;

Individual	Title	IOI Support Program(s)
		IV Therapy: An Important Option for
		Your Patients;
		Payer Relationship Management;
		Practice Compliance for Remicade;
_		Quality of Care in the Infusion Suite
Tom Nyairo	Associate Product Manager, Care Delivery	Significantly involved in the decision
	Strategies (Aug. 2004-Nov. 2006)	to stop providing any IOI Support
		Services between August 2004 and
		November 2006.
		Programs that were discontinued
		between August 2004 and November
		2006 included:
Textoody in the part of the second		Infusion Suite Designer
Bill Pillat	Product Manager, Site of Care (May 2017–	Significantly involved in the decision
	Aug. 2019)	to stop providing any IOI Support
		Services between May 2017 and
		August 2019.
		Programs that were discontinued
		between May 2017 and August 2019
		included:
		Billing and Coding for Infusions;
		Emerging Trends in Healthcare;
		Infusion Services Review (iBiz);
		Infusion Suite Scheduling and Staffing;
		IV Therapy: An Important Option for
		Your Patients;
		Payer Relationship Management;
		Quality of Care in the Infusion Suite
Faiz Sadeq	Product Manager, Alternative Site of Care	Significantly involved in the decision
	(Dec. 2010–July 2011);	to stop providing any IOI Support
	Product Manager, Site of Care/Hospital	Services between December 2010 and
	(July 2011–Dec. 2012)	December 2012.
		Programs that were discontinued
		between December 2010 and
		December 2012 included:
		Managing Biologics in the Physician
		Office;
		Successful Infusion Site Management
		for Gastroenterology;

Individual	Title	IOI Support Program(s)
		Successful Implementation of a New Infusion Suite for Gastroenterology
		Practices;
		Successful Implementation of a New
		Infusion Suite;
		Private Payer Contracting
		Considerations – Part 1 and Part 2
		(a/k/a Private Payer Contracting
		Considerations for Therapy With
		Remicade);
		Enhancing Efficiency and Capacity;
		Considerations for Working With a
		Specialty Pharmacy;
		Becoming an Alternative Site of Care
		for Therapy with Remicade in Your
	D : 1D : D: (1 2015	Community
Scott	Regional Business Director (Apr. 2015–	Significantly involved in the decision
Shelhamer	Feb. 2020)	to stop providing any IOI Support Services between April 2015 and
		February 2020.
Sandra	Product Manager (2002–2005)	Significantly involved in the decision
Shpilberg	1 Toddet Wanager (2002-2003)	to stop providing any IOI Support
Shphoerg		Services between 2002 and 2005.
		Services services 2002 and 2000.
		Programs that were discontinued
		between 2002 and 2005 included:
		Infusion Suite Designer
Karen Trahan	Regional Business Director (2000–2015)	Significantly involved in the decision
		to stop providing any IOI Support
3 6' 1 1	B 1 B 1 C 1 C 1 C	Services between 2000 and 2015.
Michael	Product Director, Marketing, Site of Care	Significantly involved in the decision
Wolfe	(Oct. 2010–Apr. 2012);	to stop providing any IOI Support
	Product Director, Rheumatology	Services between October 2010 and
	Marketing (Apr. 2012–Apr. 2016)	April 2016.
		Programs that were discontinued
		between October 2010 and April 2016
		included:
		Becoming an Alternative Site of Care
		for Therapy with Remicade in Your
		Community;
		Checkpoints for Infusion Center
		Optimization;

Individual	Title	IOI Support Program(s)
		Considerations for Working With a
		Specialty Pharmacy;
		Enhancing Efficiency and Capacity;
		Infusion Optimization Modeler (IOM);
		Infusion Therapy Services Provided in
		Converted ASC Space (a/k/a Infusion
		Services and Ambulatory Surgical
		Centers (ASCs) – Planning
		Considerations;
		ASC Space Reclassification for
		Infusion Therapy);
		Practice Compliance for Remicade;
		Private Payer Contracting
		Considerations – Part 1 and Part 2
		(a/k/a Private Payer Contracting
		Considerations for Therapy With
		Remicade);
		Setting Up In-Office Infusions of
		Remicade: Informational Resources;
		Successful Implementation of a New
		Infusion Suite;
		Successful Implementation of a New
		Infusion Suite for Gastroenterology
		Practices;
		Successful Infusion Site Management
		for Gastroenterology (a/k/a Successful
		Infusion Suite Management for
		Gastroenterology)
David Wright	Group Product Director, Stakeholder (Feb.	Significantly involved in the decision
David Wilght	2013–Dec. 2014);	to stop providing any IOI Support
	Group Product Director, Rheumatology,	Services between February 2013 and
	IV/IOI (Dec. 2014–Mar. 2016)	March 2016.
	11/101 (Bec. 2014 Mai. 2010)	March 2010.
		Programs that were discontinued
		between February 2013 and March
		2016 included:
		TO A STATE OF THE PARTY OF THE
		Checkpoints for Infusion Center
		Optimization;
		Infusion Optimization Modeler (IOM);
		Infusion Therapy Services Provided in
		Converted ASC Space (a/k/a Infusion
		Services and Ambulatory Surgical
		Centers (ASCs) – Planning
		Considerations;

Individual	Title	IOI Support Program(s)
		ASC Space Reclassification for
		Infusion Therapy);
		Practice Compliance for Remicade;
		Setting Up In-Office Infusions of
		Remicade: Informational Resources
Michael	Director, Public Payer Policy, Strategy &	Significantly involved in the decision
Ziskind	Marketing (Centocor) (1998-2006)	to stop providing any IOI Support
		Services between 1998 and 2006.
		Programs that were discontinued
		between 1998 and 2006 included:
		Infusion Suite Designer

<u>TOPIC NO. 3:</u> The persons (excluding employees acting in the capacity of an Area Business Specialist or Regional Business Manager) that had significant involvement in assessing, evaluating, and/or analyzing customer demand for and the value customers derived from any or all of the IOI Support Services (including the specific IOI Support Service(s) in which each such person had the significant involvement, the approximate date(s) of the significant involvement, the person's last known address and phone number, and, if an individual, the position they held at the time of their significant involvement).

Relevant time period: From October 28, 2006 to until February 19, 2016.

RESPONSE TO TOPIC NO. 3:

Janssen objects to this Topic to the extent that it is overly broad and unduly burdensome and not proportional to the needs of the case, and thus exceeds the scope of discoverable matters under Federal Rule of Civil Procedure 26.

Janssen further objects to this Topic to the extent it is cumulative and seeks information that is duplicative of what is contained in the documents or written discovery that Janssen has already produced.

Janssen further objects that this Topic on the ground that it is vague, ambiguous, and overbroad and does not describe with reasonable particularity the matters on which examination is requested as required by Fed. R. Civ. P. 30(b)(6). For example, Janssen objects to the Topic's use of the terms "significant involvement," "customer demand," and "value," as those are not

defined terms. "Significant involvement," in particular, is an inherently subjective term, and reasonable minds could differ as to whether a particular person was "significantly" involved in any given activity. This response is based on Janssen's good faith understanding, and it seeks to address Relator's concerns about providing a response that is over-inclusive or under-inclusive.

Further, pursuant to the Court's March 9, 2023 order, Janssen's response will "only be with regard to the services at issue in this case, not all services." *See* ECF No. 375 at 18.

Accordingly, Janssen's response to Topic No. 3 will be with respect to the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2. *See* Relator's 3d Supplemental Objections and Responses to Janssen's Rog. No. 2 (January 27, 2023).

Subject to and without waiving the foregoing objections, Janssen responds as follows:

The Site of Care team, with input from the ABSs, is responsible for determining the educational materials available for use by the ABSs. *See, e.g.*, JANSSENBIO-011-00002691; JANSSENBIO-011-00002730; M. Wolfe Dep. Tr. at 74:16-75:22, 256:21-257:4. The determination as to which educational materials would be delivered by the ABSs was based upon the ABSs continued usage of educational materials (*see* JANSSENBIO-011-00002691; JANSSENBIO-011-00002730), redundancy of content across educational materials (*see* JANSSENBIO-014-00001575), and the educational needs of practices that were infusing Remicade and Simponi ARIA, as identified by ABSs (*see* M. Wolfe Dep. Tr. at 74:16-75:22). The individuals responsible for determining the educational materials available for use by the ABSs are identified in response to Topic 1.

Based on its investigation to date, Janssen is unaware of anyone assessing, evaluating, or analyzing "customer demand for" or the "value customers derived from" the IOI Support Services identified by Relator in response to Interrogatory No. 2.

<u>TOPIC NO. 4:</u> The persons (excluding the PRC) that had significant involvement in reviewing, assessing, and/or advising you regarding the legality or illegality of providing any or all of the IOI Support Services (including the specific IOI Support Service(s) in which each such person had the significant involvement, the approximate date(s) of the significant involvement, the person's last known address and phone number, and, if an individual, the position they held at the time of their significant involvement).

<u>Relevant time period:</u> From when the strategy of providing the IOI Support Services was devised until February 19, 2020.

RESPONSE TO TOPIC NO. 4:

Janssen objects to this Topic to the extent that it is overly broad and unduly burdensome and not proportional to the needs of the case, and thus exceeds the scope of discoverable matters under Federal Rule of Civil Procedure 26.

Janssen further objects to this Topic to the extent it is cumulative and seeks information that is duplicative of what is contained in the documents or written discovery that Janssen has already produced.

Janssen further objects that this Topic on the ground that it is vague, ambiguous, and overbroad and does not describe with reasonable particularity the matters on which examination is requested as required by Fed. R. Civ. P. 30(b)(6). For example, Janssen objects to the Topic's use of the term "significant involvement," as that is not a defined term. "Significant involvement," in particular, is an inherently subjective term, and reasonable minds could differ as to whether a particular person was "significantly" involved in any given activity. This response is based on Janssen's good faith understanding, and it seeks to address Relator's concerns about providing a response that is over-inclusive or under-inclusive.

Further, pursuant to the Court's March 9, 2023 order, Janssen's response will "only be with regard to the services at issue in this case, not all services." *See* ECF No. 375 at 18.

Accordingly, Janssen's response to Topic No. 4 will be with respect to the IOI Support Services

identified by Relator in response to Janssen's Interrogatory No. 2. *See* Relator's 3d Supplemental Objections and Responses to Janssen's Rog. No. 2 (January 27, 2023).

As previously stated, Janssen has not asserted and does not intend to assert the advice of counsel defense. In answering this Topic, Janssen is not disclosing or intending to imply anything about what legal advice may or may not have been provided. For avoidance of doubt, by providing the requested factual information regarding the identities of the individuals who provided privileged legal advice, Janssen is not asserting the advice of counsel defense and has not waived, and has no intention of waiving, any applicable privilege.

Subject to and without waiving the foregoing objections, Janssen responds as follows:

The following list encompasses the Janssen individuals significantly involved in reviewing, assessing, and/or advising Janssen regarding the legality or illegality of providing the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2:

Individual	Title	IOI Support Program
Chris Guiton	Assistant General Counsel (July 2007–Mar. 2019);	General legal advice regarding SOC educational materials, including IOI
	Senior Director, US State Government Affairs (Mar. 2019–present)	Support Services, between July 2007 and March 2019.
		Listed as the Legal Reviewer on relevant PRC forms, including for at
		least the following IOI Support Services:
		Infusion Services Review (iBiz); Raising the Infusion Suite Experience (RISE)
Kathleen Hamill	Assistant General Counsel, Centocor Board (Apr. 2000–2011);	General legal advice regarding SOC educational materials, including IOI
	Assistant General Counsel, Regulatory (2011–present)	Support Services, between April 2000 and present.
		Listed as the Legal Reviewer on relevant PRC forms, including for at

Individual	Title	IOI Support Program
		least the following IOI Support
		Services:
		Checkpoints for Infusion Center
		Optimization;
		Managing Biologics in the Physician
		Office;
		Private Payer Contracting
		Considerations;
		Successful Infusion Suite
F., 11, T.,	A - i - t t C 1 C 1 (S t	Management for Gastroenterology
Freddy Jimenez	Assistant General Counsel (Sept.	General legal advice regarding SOC
	1999–Jan. 2016)	educational materials, including IOI
		Support Services, between 1999 and
		2016.
		Listed as the Legal Reviewer on
		relevant PRC forms, including for at
		least the following IOI Support
		Services:
		Services.
		Becoming an Alternative Site of Care;
		Considerations for Proactive Practice
		Management; Emerging Trends in
		Healthcare;
		Infusion Optimization Modeler
		(IOM);
		Practice Compliance for Remicade;
		Private Payer Contracting
		Considerations for Remicade;
		Remicade Account Review;
		Setting Up In-Office Infusions of
		Remicade
Michael McCulley	Assistant General Counsel (Apr.	General legal advice regarding SOC
111 33	1982–Dec. 2012)	educational materials, including IOI
		Support Services, between 2000 and
		2012.
Diedre Meehan	Assistant General Counsel, Regulatory	General legal advice regarding SOC
	(Aug. 2007–Dec. 2020)	educational materials, including IOI
		Support Services, between August
Control of the Contro		2007 and December 2020.
Daryl Todd	Senior Counsel, Regulatory (Apr.	General legal advice regarding SOC
	2010–June 2015);	educational materials, including IOI
	Assistant General Counsel, Regulatory	Support Services, between 2010 and
	(June 2015–present)	present.

Individual	Title	IOI Support Program
John Vaughan	Senior Counsel (2007–June 2012)	General legal advice regarding SOC
		educational materials, including IOI
		Support Services, between 2007 and
		June 2012.
		Listed as the Legal Reviewer on
		relevant PRC forms, including for at
		least the following IOI Support
		Services:
		Emerging Trends in Healthcare;
		Enhancing Infusion Efficiency;
		Infusion Optimization Modeler
		(IOM);
		Infusion Therapy Services Provided in
		Converted ASC Space;
		IV Therapy: An Important Option for
		Your Patients (a/k/a Why IV?);
		Managing Biologics in the Physician
		Office;
		Private Payer Contracting
		Considerations; Successful
		Implementation of a New Infusion
		Suite;
		Successful Implementation of a New
		Infusion Suite for Gastroenterology
		Practices; Successful Infusion Suite
		Management for Gastroenterology

Additionally, Janssen also retained outside counsel from Reed Smith, Hogan & Hartson, and Akin Gump.

TOPIC NO. 5: The persons who analyzed, reviewed, and/or advised whether you should or should not seek an advisory opinion from OIG concerning any of the IOI Support Services (including the specific IOI Support Service(s) addressed or covered by the analysis/analyses, review(s), or advice each such person performed or provided, the approximate date(s), the person's last known address and phone number, and, if an individual, the position held at the time the analysis/analyses, review(s), or advice was/were performed or provided).

<u>Relevant time period</u>: From when the strategy of providing the IOI Support Services was devised until February 19, 2020.

RESPONSE TO TOPIC NO. 5:

Janssen objects to this Topic to the extent that it is overly broad and unduly burdensome and not proportional to the needs of the case, and thus exceeds the scope of discoverable matters under Federal Rule of Civil Procedure 26.

Further, pursuant to the Court's March 9, 2023 order, Janssen's response will "only be with regard to the services at issue in this case, not all services." *See* ECF No. 375 at 18.

Accordingly, Janssen's response to Topic No. 5 will be with respect to the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2. *See* Relator's 3d Supplemental Objections and Responses to Janssen's Rog. No. 2 (January 27, 2023).

Subject to and without waiving the foregoing objections, Janssen responds as follows:

There are no persons who analyzed, reviewed, and/or advised whether Janssen should or should not seek an advisory opinion from OIG concerning any of the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2.

TOPIC NO. 6: The persons (excluding the PRC) that had significant involvement in reviewing, assessing, and/or advising you whether providing any or all of the IOI Support Services complied with your compliance policies (including the specific IOI Support Service(s) addressed or covered by the review(s), assessment(s), or advice each such person performed or provided, the approximate date(s) of the significant involvement, the person's last known address and phone number, and, if an individual, the position they held at the time of their significant involvement).

<u>Relevant time period:</u> From when the strategy of providing the IOI Support Services was devised until February 19, 2020.

RESPONSE TO TOPIC NO. 6:

Janssen objects to this Topic to the extent that it is overly broad and unduly burdensome and not proportional to the needs of the case, and thus exceeds the scope of discoverable matters under Federal Rule of Civil Procedure 26.

Janssen further objects to this Topic to the extent it is cumulative and seeks information that is duplicative of what is contained in the documents or written discovery that Janssen has already produced.

Further, pursuant to the Court's March 9, 2023 order, Janssen's response will "only be with regard to the services at issue in this case, not all services." *See* ECF No. 375 at 18.

Accordingly, Janssen's response to Topic No. 6 will be with respect to the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2. *See* Relator's 3d Supplemental Objections and Responses to Janssen's Rog. No. 2 (January 27, 2023).

Janssen further objects that this Topic on the ground that it is vague, ambiguous, and overbroad and does not describe with reasonable particularity the matters on which examination is requested as required by Fed. R. Civ. P. 30(b)(6). For example, Janssen objects to the Topic's use of the term "significant involvement," as that is not a defined term. "Significant involvement," in particular, is an inherently subjective term, and reasonable minds could differ as to whether a particular person was "significantly" involved in any given activity. This response is based on Janssen's good faith understanding, and it seeks to address Relator's concerns about providing a response that is over-inclusive or under-inclusive.

As previously stated, Janssen has not asserted and does not intend to assert the advice of counsel defense. In answering this Topic, Janssen is not disclosing or intending to imply anything about what legal advice may or may not have been provided. For avoidance of doubt, by providing the requested factual information regarding the identities of the individuals who provided privileged legal advice, Janssen is not asserting the advice of counsel defense and has not waived, and has no intention of waiving, any applicable privilege.

Subject to and without waiving the foregoing objections, Janssen responds as follows:

The following list encompasses the individuals significantly involved in reviewing, assessing, and/or advising Janssen whether providing the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2 complied with Janssen's compliance policies:

Individual	Title	IOI Support Program
Michele Blades	Health Care Compliance Officer (Mar. 2017–present)	General compliance advice regarding SOC educational materials, including IOI Support Services, between March 2017 and February 2020.
		Listed as the HCC Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Infusion Services Review (iBiz); Inventory and Supply Management; Infusion Practice Management Series - Part 1; Infusion Practice Management Series - Part 2; Billing and Coding for Infusions; Quality of Care in the Infusion Suite; In-Office Drug Procurement Models; Infusion Suite Scheduling and Staffing; ICD-10; Specialty Drug Market Dynamics:
Thomas Cornely	Health Care Compliance Officer, Immunology (Apr. 2013–2016)	Implications for Infusions General compliance advice regarding SOC educational materials, including IOI Support Services, between 2013 and 2016. Listed as the HCC Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Billing and Coding for Infusions; Infusion Services Review (iBiz);

Individual	Title	IOI Support Program
		Infusion Suite Scheduling and Staffing; Infusion Practice Management Series - Part 1; Infusion Practice Management Series - Part 2; In-Office Drug Procurement Models; Billing and Coding for Infusions; Payer Relationship Management; Inventory and Supply Management; ICD-10
John "Chip" Franz	Health Care Compliance Officer (Oct. 2010–Aug. 2014)	General compliance advice regarding SOC educational materials, including IOI Support Services, between 2010 and 2014. Listed as the HCC Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Emerging Trends in Healthcare; Infusion Services Review (iBiz); IV Therapy: An Important Option for Your Patients (a/k/a Why IV?); Raising the Infusion Suite Experience (RISE)
Gina Giordano	Health Care Compliance Officer (June 2009–Oct. 2015)	General compliance advice regarding SOC educational materials, including IOI Support Services, between 2009 and 2015. Listed as the HCC Reviewer on relevant PRC forms, including for at least the following IOI Support Services: Considerations for Working with a Specialty Pharmacy
Edmund Greenidge	Director, Health Care Compliance (2005–Oct. 2010)	General compliance advice regarding SOC educational materials, including IOI Support Services, between 2005 and 2010.

Individual	Title	IOI Support Program
		Listed as the HCC Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Becoming an Alternative Site of Care; Enhancing Patient Care and Access; Considerations for Proactive Practice Management; Infusion Therapy Services Provided in Converted ASC Space; Managing Biologics in the Physician Office; Setting Up In-Office Infusions of
Chris Guiton	Assistant General Counsel (July 2007–Mar. 2019); Senior Director, US State Government Affairs (Mar. 2019–present)	Remicade General legal advice regarding SOC educational materials, including IOI Support Services, between July 2007 and March 2019.
		Listed as the Legal Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Infusion Services Review (iBiz); Raising the Infusion Suite Experience (RISE)
Kathleen Hamill	Assistant General Counsel, Centocor Board (Apr. 2000–2011); Assistant General Counsel, Regulatory (2011–present)	General legal advice regarding SOC educational materials, including IOI Support Services, between April 2000 and present.
		Listed as the Legal Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Checkpoints for Infusion Center Optimization; Managing Biologics in the Physician Office; Private Payer Contracting Considerations;

Individual	Title	IOI Support Program
		Successful Infusion Suite
		Management for Gastroenterology
Freddy Jimenez	Assistant General Counsel (Sept. 1999–Jan. 2016)	General legal advice regarding SOC educational materials, including IOI Support Services, between 1999 and 2016.
		Listed as the Legal Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Becoming an Alternative Site of Care; Considerations for Proactive Practice Management; Emerging Trends in Healthcare; Infusion Optimization Modeler (IOM); Practice Compliance for Remicade; Private Payer Contracting Considerations for Remicade; Remicade Account Review; Setting Up In-Office Infusions of Remicade
Roger Kung	Health Care Compliance Officer (2011–2012)	General compliance advice regarding SOC educational materials, including IOI Support Services, between 2011 and 2012. Listed as the HCC Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Checkpoints for Infusion Center Optimization; Managing Biologics in the Physician Office; Private Payer Contracting Considerations
Michael McCulley	Assistant General Counsel (Apr. 1982–Dec. 2012)	General legal advice regarding SOC educational materials, including IOI Support Services, between 2000 and 2012.

Individual	Title	General legal advice regarding SOC educational materials, including IOI Support Services, between August 2007 and December 2020.	
Diedre Meehan	Assistant General Counsel, Regulatory (Aug. 2007–Dec. 2020)		
Erin Parsons	Health Care Compliance Officer (Jan. 2015–Nov. 2019)	General compliance advice regarding SOC educational materials, including IOI Support Services, between 2015 and 2019.	
		Listed as the HCC Reviewer on relevant PRC forms, including for at least the following IOI Support Services:	
		ICD-10; Specialty Drug Market Dynamics: Implications for Infusions	
Maripat Rhood	Manager, Health Care Compliance (May 2001–July 2007)	General compliance advice regarding SOC educational materials, including IOI Support Services, between May 2001 and July 2007.	
Michael Schoeck	Director, HCC and Privacy (Oct. 2002–Aug. 2008); Director, HCC (Aug. 2008–Dec. 2010)	General compliance advice regarding SOC educational materials, including IOI Support Services, between 2002 and 2010.	
		Listed as the HCC Reviewer on relevant PRC forms, including for at least the following IOI Support Services:	
		Private Payer Contracting Considerations for Remicade; Remicade Account Review	
Daryl Todd	Senior Counsel, Regulatory (Apr. 2010–June 2015); Assistant General Counsel, Regulatory (June 2015–present)	General legal advice regarding SOC educational materials, including IOI Support Services, between 2010 and present.	
John Vaughan	Senior Counsel (2007–June 2012)	General legal advice regarding SOC educational materials, including IOI Support Services, between 2007 and June 2012.	
		Listed as the Legal Reviewer on relevant PRC forms, including for at	

Individual	Title	IOI Support Program
		least the following IOI Support
		Services:
		Emerging Trends in Healthcare;
		Enhancing Infusion Efficiency;
		Infusion Optimization Modeler
		(IOM);
		Infusion Therapy Services Provided
		in Converted ASC Space;
		IV Therapy: An Important Option for
		Your Patients (a/k/a Why IV?);
		Managing Biologics in the Physician
		Office;
		Private Payer Contracting
		Considerations; Successful
		Implementation of a New Infusion
		Suite;
		Successful Implementation of a New
		Infusion Suite for Gastroenterology
		Practices; Successful Infusion Suite
4 1 777 1	H H C C I' OW OI	Management for Gastroenterology
Angela Wood	Health Care Compliance Officer (Nov.	General compliance advice regarding
	2010–June 2014)	SOC educational materials, including
		IOI Support Services, between 2010 and 2014.
		and 2014.
		Listed as the HCC Reviewer on
		relevant PRC forms, including for at
		least the following IOI Support
		Services:
		Services.
		Emerging Trends in Healthcare
Chris Zalesky	Director/Executive Director,	General compliance advice regarding
	Marketing Compliance (Centocor)	SOC educational materials, including
	(April 1998–Oct. 1999);	IOI Support Services, between 1998
	Director, Regulatory Affairs (Janssen)	and 2010.
	(Oct. 1999–Mar. 2002);	
	Senior Director, Regulatory Affairs	
	(Janssen) (Mar. 2002–May 2004);	
	Executive Director, World Wide	
	Office of Health Care Compliance and	
	Policy (June 2004–Mar. 2010);	
	Vice President, Global Policy &	
	Guidance (2010–2016)	

TOPIC NO. 7: The persons that performed the analyses or reviews and/or provided the opinions or advice concerning the HHS or OIG guidance, OIG advisory opinions, Anti-Kickback Statute, False Claims Act, federal regulations, and/or judicial decisions referenced in your responses to interrogatories 15 (amended) and 20-22 that you considered, relied upon, and/or adopted in reaching the asserted belief at the time you were providing the IOI Support Services that you were acting lawfully (including the specific IOI Support Service(s) addressed or covered by the analysis/analyses, review(s), opinion(s), or advice each such person performed or provided, the approximate date(s), the person's last known address and phone number, and, if an individual, the position held at the time the analysis/analyses, review(s), opinion(s), or advice was/were performed or provided).

<u>Relevant time period:</u> From when the strategy of providing the IOI Support Services was devised until February 19, 2020.

RESPONSE TO TOPIC NO. 7:

Janssen objects to this Topic to the extent that it is overly broad and unduly burdensome and not proportional to the needs of the case, and thus exceeds the scope of discoverable matters under Federal Rule of Civil Procedure 26.

Janssen further objects to this Topic to the extent it is cumulative and seeks information that is duplicative of what is contained in the documents or written discovery that Janssen has already produced.

Janssen further objects to this Topic to the extent it mischaracterizes Janssen's response to Interrogatory Nos. 15 and 20–22.

Further, pursuant to the Court's March 9, 2023 order, Janssen's response will "only be with regard to the services at issue in this case, not all services." *See* ECF No. 375 at 18.

Accordingly, Janssen's response to Topic No. 7 will be with respect to the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2. *See* Relator's 3d Supplemental Objections and Responses to Janssen's Rog. No. 2 (January 27, 2023).

As previously stated, Janssen has not asserted and does not intend to assert the advice of counsel defense. In answering this Topic, Janssen is not disclosing or intending to imply anything about what legal advice may or may not have been provided. For avoidance of doubt,

by providing the requested factual information regarding the identities of the individuals who provided privileged legal advice, Janssen is not asserting the advice of counsel defense and has not waived, and has no intention of waiving, any applicable privilege.

Subject to and without waiving the foregoing objections, Janssen responds as follows:

The following list encompasses the individuals significantly involved in the analyses or reviews and/or provided the opinions or advice concerning the legality of the IOI Support Services identified by Relator in response to Janssen's Interrogatory No. 2:

Individual	Title	IOI Support Program
Chris Guiton	Assistant General Counsel (July 2007–Mar. 2019); Senior Director, US State Government Affairs (Mar. 2019–present)	General legal advice regarding SOC educational materials, including IOI Support Services, between July 2007 and March 2019. Listed as the Legal Reviewer on
		relevant PRC forms, including for at least the following IOI Support Services:
		Infusion Services Review (iBiz); Raising the Infusion Suite Experience (RISE)
Kathleen Hamill	Assistant General Counsel, Centocor Board (Apr. 2000–2011); Assistant General Counsel, Regulatory (2011–present)	General legal advice regarding SOC educational materials, including IOI Support Services, between April 2000 and present.
		Listed as the Legal Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Checkpoints for Infusion Center Optimization; Managing Biologics in the Physician Office;
		Private Payer Contracting Considerations; Successful Infusion Suite Management for Gastroenterology

Individual	Title	IOI Support Program
Freddy Jimenez	Assistant General Counsel (Sept. 1999–Jan. 2016)	General legal advice regarding SOC educational materials, including IOI Support Services, between 1999 and 2016.
		Listed as the Legal Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Becoming an Alternative Site of Care; Considerations for Proactive Practice Management; Emerging Trends in Healthcare; Infusion Optimization Modeler (IOM); Practice Compliance for Remicade; Private Payer Contracting Considerations for Remicade; Remicade Account Review; Setting Up In-Office Infusions of
Mike McCulley	Assistant General Counsel (Apr. 1982–Dec. 2012)	Remicade General legal advice regarding SOC educational materials, including IOI Support Services, between 2000 and 2012.
Diedre Meehan	Assistant General Counsel, Regulatory (Aug. 2007–Dec. 2020)	General legal advice regarding SOC educational materials, including IOI Support Services, between August 2007 and December 2020.
Daryl Todd	Senior Counsel, Regulatory (Apr. 2010–June 2015); Assistant General Counsel, Regulatory (June 2015–present)	General legal advice regarding SOC educational materials, including IOI Support Services, between 2010 and present.
John Vaughan	Senior Counsel (2007–June 2012)	General legal advice regarding SOC educational materials, including IOI Support Services, between 2007 and June 2012.
		Listed as the Legal Reviewer on relevant PRC forms, including for at least the following IOI Support Services:
		Emerging Trends in Healthcare;

Individual	Title	IOI Support Program
		Enhancing Infusion Efficiency;
		Infusion Optimization Modeler
		(IOM);
		Infusion Therapy Services Provided in
		Converted ASC Space;
		IV Therapy: An Important Option for
		Your Patients (a/k/a Why IV?);
		Managing Biologics in the Physician
		Office;
		Private Payer Contracting
		Considerations; Successful
		Implementation of a New Infusion
		Suite;
		Successful Implementation of a New
		Infusion Suite for Gastroenterology
		Practices; Successful Infusion Suite
		Management for Gastroenterology

Additionally, Janssen retained outside counsel from Reed Smith, Hogan & Hartson, and Akin Gump.

TOPIC NO. 8: The persons that performed the analysis(es) or review(s) and/or provided the opinions or advice concerning the meaning, implication, or significance of the United States' investigation or inquiry referenced in your response to interrogatories 15 (amended) and 20-22 that you considered, relied upon, and/or adopted in reaching the asserted belief at the time you were providing the IOI Support Services that you were acting lawfully (including the specific IOI Support Service(s) addressed or covered by the analysis/analyses, review(s), opinion(s), or advice each such person performed or provided, the approximate date(s), the person's last known address and phone number, and, if an individual, the position held at the time the analysis/analyses, review(s), opinion(s), or advice was/were performed or provided).

Relevant time period: From 2003 until February 19, 2020.

RESPONSE TO TOPIC NO. 8:

Janssen objects to this Topic to the extent that it is overly broad and unduly burdensome and not proportional to the needs of the case, and thus exceeds the scope of discoverable matters under Federal Rule of Civil Procedure 26.

Janssen further objects to this Topic to the extent it is cumulative and seeks information that is duplicative of what is contained in the documents or written discovery that Janssen has already produced.

Janssen also objects to any implication that the reasonableness of its interpretation of the statutes and regulations at issue in the SAC can only be supported by contemporaneous, affirmative analyses that were performed at a particular time.

Janssen further objects to this Topic to the extent it mischaracterizes Janssen's response to Interrogatory Nos. 15 and 20–22.

As previously stated, Janssen has not asserted and does not intend to assert the advice of counsel defense. In answering this Topic, Janssen is not disclosing or intending to imply anything about what legal advice may or may not have been provided. For avoidance of doubt, by providing the requested factual information regarding the identities of the individuals who provided privileged legal advice, Janssen is not asserting the advice of counsel defense and has not waived, and has no intention of waiving, any applicable privilege.

Subject to and without waiving the foregoing objections, Janssen responds as follows:

The following list encompasses the individuals significantly involved in the United States' investigation referenced in Janssen's response to Relator's Interrogatory Nos. 15 (amended) and 20–22, beginning in 2003:

Individual	Role/Dates
Joseph Braunreuther	General Attorney (Aug. 2001–June 2006); General Counsel, Pharmaceuticals Group (July 2006–Dec. 2008);
	Associate General Counsel, Litigation (Jan. 2009–Dec. 2011); Deputy General Counsel (Jan. 2012–Dec. 2021)
Freddy Jimenez	Assistant General Counsel (Sept. 1999–Jan. 2016)

Additionally, Janssen retained outside counsel from Reed Smith.

VERIFICATION

I am the Vice President, Immunology Portfolio Strategy at Janssen Biotech, Inc. I am authorized to make this verification on behalf of Janssen Biotech, Inc. I have read Defendant's foregoing objections and responses to Plaintiff-Relator Julie Long's 30(b)(6) Deposition Notice served on November 4, 2022. I swear under penalty of perjury that the information provided is true and accurate to the best of my knowledge, information and belief.

Executed on March 23, 2023.

Brian Smith

AS TO OBJECTIONS AND RESPONSES:

Dated: March 23, 2023 s/Jason C. Raofield

Jason C. Raofield (BBO No. 641744)
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CERTIFICATE OF SERVICE

The undersigned hereby certifies that a true and correct copy of the foregoing document has been served by electronic mail on March 23, 2023, to the following counsel of record:

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